INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10557511	
Filing Date		2005-11-21	
First Named Inventor	Jee I	ee Ho KIM	
Art Unit		1795	
Examiner Name	NYA		
Attorney Docket Number		LEE-0042	

CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patient office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(eV1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign pattern office in a counterpart foreign application, and, to the knowledge of the person signing the certification are making reasonable inquiry, no item of information contained in the information disclosure statement was known to any inclodual designated in 70 CPR 1.56(c) more than three morning port to the filling of the information disclosure any inclodual designated in 70 CPR 1.56(c) more than three morning port to the filling of the information disclosure and the contract of the contr

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

Dana A. Gronbeck

□ None

Name/Print

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

form of the signature.			
Signature	/Dana A. Gronbeck/	Date (YYYY-MM-DD)	2009-06-11

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is folling and by the USPTO to process) an application. Confidentially is governed by \$5.0 S.C. 12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Pattert and Trademark Office, U.S. Operament of Commence, P.O. Boat 1450, Alexandria, V.A.2311-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.A.2311-1450.

Registration Number

55226

Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is SU S.C. 2(b)(2); (2) furnishing of the information solicited is civilarity; and (5) the primoral purpose for which the information is used by the U.S. Patient and Trademan KOTIes is to information, the U.S. Patient and Trademan KOTIes may not be able to process and/or examine your submission, which may result in farmination of proceedings or abandoment of the explication or experients of the patient.

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement necodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, cursuant to 5 U.S.C. 552(m).
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 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher dissigne, during an inspection of records conducted by GSA a part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 12(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application poen to public insepticines or an insuce patent.
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